

**Report of CDC IRB “ ”**  
**CDC Protocol # \_\_\_\_\_, “ ”**  
**April 23, 2001**

**General Comments and IRB Actions**

The convened board of IRB “ ” reviewed your request (for new protocol approval/to continue/to amend protocol # \_\_\_\_\_ . We have determined that the study (still) involves (greater than/no more than) minimal risk to subjects. Upon receipt of satisfactory responses to the following issues and concerns, and upon receipt of a clean copy/ies of the revised protocol/consent form(s)/assent form(s)/questionnaire(s), the IRB will approve (continuation of your/your amended/ your new) protocol.

**Protocol Issues**

Response Required, Action Required

- 1)
- 2)
- 3)

Response Required, Action Optional

- 1)
- 2)
- 3)

Of Note - No Response or Action Required

**Consent Form Issues**

Response Required, Action Required

- 1)
- 2)
- 3)

Response Required, Action Optional

- 1)
- 2)

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3)

*Of Note - No Response or Action Required*

**Addenda Issues (Questionnaires, brochures, posters, etc.)**

*Response Required, Action Required*

- 1)
- 2)
- 3)

*Response Required, Action Optional*

- 1)
- 2)
- 3)

*Of Note - No Response or Action Required*

**End of Report**